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### **Original** Article



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# The association of glycemic index and glycemic load with elevated blood pressure in Iranian women

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#### Abstract

*Introduction:* Dietary intake is a risk factor related to elevated blood pressure (EBP). Few studies have investigated an association of dietary glycemic index (GI) and glycemic load (GL) with the EBP. The aim of the current study was to examine the association of dietary GI and GL with the EBP among a group of healthy women.

*Methods:* This population-based cross-sectional study was conducted on 306 healthy women. Dietary GI and GL were measured using a validated semi-quantitative food frequency questionnaire (FFQ). Blood pressure (BP) was measured twice by a mercury sphygmomanometer from the right arm. Anthropometric measurements were also assessed according to the standard protocols.

**Results:** Before controlling for potential confounders, no significant association was seen between dietary GI/GL and SBP/DBP. Also after controlling for potential confounders, the associations did not change between dietary GI and SBP (odds ratio [OR]: 0.96; 95% CI: 0.42-2.17, P=0.87), between GI and DBP (OR: 0.72; 95% CI: 0.35-1.45, P=0.37), as well as between GL and SBP (OR: 1.04; 95% CI: 0.43-2.49, P=1.00) and between GL and DBP (OR: 1.20; 95% CI: 0.56-2.00, P=0.61). In a stratified analysis by obesity and overweight, differences between tertiles of GI were not significant (OR: 0.75; 95% CI: 0.42-1.31, P=0.31), even after adjustment for the potential confounders (OR: 1.54; 95% CI: 0.70-3.40, P=0.26).

*Conclusion:* This study did not show a significant association between dietary GI/GL and the risk of high SBP/DBP. In addition, no significant association was found between dietary GI/GL and odds of overweight or obesity in adult women.

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#### Introduction

Elevated blood pressure (EBP), based on adult treatment panel III, is defined as a systolic blood pressure (SBP)  $\geq$ 130 mmHg and/or diastolic blood pressure (DBP)  $\geq$ 85 mm Hg.<sup>1</sup> Prevalence of hypertension has gently increased in the 2 last decades. Hypertension has been estimated to cause annually 7.5 million deaths worldwide.<sup>2,3</sup> Unfortunately, prevalence of hypertension is high among Iranians.<sup>4,5</sup> Hypertension causes different micro- and macro- vascular complications and increases all-cause and cardiovascular disease mortality.<sup>6</sup> As the high complications due to EBP, its prevention is a public health priority, worldwide.

Several risk factors including genetic, environmental, lifestyle and psychosocial factors as well as dietary intake have been linked to the risk of EBP.<sup>7,8</sup> Among them, dietary

intake is one of the main environmental risk factors related to EBP.<sup>8,9</sup> For instance, intakes of sodium, potassium, calcium, magnesium, iron, phosphorus or a combination of these modalities lower blood pressure (BP).<sup>9-12</sup> In addition, dietary intake of carbohydrates has also found to be in relation with BP.<sup>10,11,13</sup> Furthermore, some studies have found no association between dietary intake of some carbohydrates and BP and showed that increased intake of dietary carbohydrate could not elevate BP,<sup>10</sup> this association was direct in some others which demonstrated diets high in carbohydrate are associated with slightly higher BP.<sup>14</sup> It seems that the association strongly depends on amount and type of the ingested carbohydrates. glycemic index (GI) is an indicator of dietary carbohydrate containing food to

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increase post-prandial blood glucose.<sup>10,15</sup> Previous studies have shown that adherence to a low GI diet may reduce BP.<sup>16,17</sup> Likewise, other studies that demonstrated excessive dietary intake of carbohydrates, particularly from high GI carbohydrates, have found to increase BP.<sup>9,20</sup> In contrast, some other studies reported no association between dietary GI and SBP/DBP.<sup>19,21</sup>

Besides to dietary GI, glycemic load (GL) is as a measure of carbohydrate quality and quantity, represents both the GI and amount of the ingested carbohydrate.<sup>22</sup> In this regard, some studies have shown a significant association between consumption of a low GL diet and decreased BP,<sup>11</sup> while others failed to find any association.<sup>23,24</sup> It should be noted that higher post-prandial glycemic response due to consumption of high GI/GL carbohydrates causes hyperinsulinemia. Hyperinsulinemia has suggested increasing sympathetic nervous system activity, which enhance heart rate, cardiac output, vascular resistance, sodium retention and thus BP.<sup>14</sup>

Despite previous studies investigated the association of GI/GL with BP in western countries; few studies have investigated this association among Iranian population, especially among healthy women. High consumption of carbohydrates including white bread and white rice, which are mainly high in GI/GL,<sup>25</sup> is common among Iranians. We conducted this study to investigate association of dietary GI/GL with EBP among healthy women referred to Tehran health centers.

#### Materials and Methods Subjects and study protocol

This is a cross-sectional study on women referring to health centers affiliated to Tehran University of Medical Sciences. Based on inclusion criteria, 306 women who were referred to health centers were selected. Inclusion criteria were satisfaction to participate in the study, being Iranian, and lack of chronic diseases such as diabetes, cardiovascular disease, hypertension, cancer, liver and kidney diseases. Those who were immigrant, pregnant or lactating were excluded. Subjects with energy intake fewer than 800 or above 4200 kcal/d were also excluded. All participants signed an informed written consent before the entrance.

#### Assessment of exposure

A validated and reliable 168-item food frequency questionnaire (FFQ) was used to assess dietary intake of participants. This semi-quantitative questionnaire consists of standard portion sizes for each food item and has been designed according to the Willett method. Participants were asked to determine the frequency of consumption of each food item during the previous year, based on serving sizes. Validity and reliability of the FFQ were determined previously.<sup>26</sup> Food intakes reported in household measures converted to grams of food per day using the Nutritionist IV software.

Total GI of participants' diet was estimated using the following formula:  $\Sigma$  (GIa \* available carbohydratea)/total available carbohydrate.27 Available carbohydrate of food items was calculated as total carbohydrate minus fiber. The total carbohydrates and fiber of 85 carbohydratecontaining food items were derived from the United State Department of Agriculture food-composition table. However, GI for 6 foods was derived from Iranian national tables.<sup>28</sup> In addition, GI values for other 62 foods were derived from the international references.<sup>29,30</sup> GI values for the rest food items which were not found neither in Iranian nor in international tables, such as some traditional desserts and sweets, were estimated based on physically and chemically similar foods.<sup>31</sup> For instance, GI value of gaz, which is chiefly made of nuts and sugar (almond or pistachios), was considered to be the same as sugar. In addition, gooshfil which mainly contains sugar and white flour, was considered the same as English muffin bread. GI values of rice and dates which have different brands, were estimated as the mean values. All derived GI values were relative to glucose as the reference food. The GIs of mixed meals were calculated based on GIs of each individual food components.<sup>27</sup> Dietary GL was estimated as (total GI \* total available carbohydrate)/100<sup>32</sup> and expressed as g/d.

#### Assessment of outcome

BP was measured by a professional clinic staff using a mercury sphygmomanometer from the right arm with appropriately sized BP cuffs. BP was measured in a sitting position and underwent measurements twice. The first measurement was taken after 5 minutes of resting. After a 15-minute rest, a second measurement was taken; the average of the 2 values is reported. In this study the description of EBP, based on adult treatment panel III, used to define a high SBP ( $\geq$ 130 mm Hg) and/or high DBP ( $\geq$  85 mm Hg).

#### Assessment of anthropometric measures

Anthropometric measurements including participants' weight, height and waist circumference were measured according to the standard protocols.<sup>33</sup> Height and weight were measured when wearing minimal clothing and without shoes using stadiometer and digital scale to the nearest of 0.1 cm and 100 g, respectively. Body mass index (BMI) was calculated as weight (kg) divided by height squared (m<sup>2</sup>). Waist circumference was measured in the middle of distance between the lowest gear and top of the iliac crest (narrowest girth), in a standing position and at the end of normal exhalation, when wearing minimum clothing to the nearest of 0.1 cm.

#### Assessment of other variables

General information including participants' name, age, gender, location and contact number were recruited using a questionnaire. Socio-economic status of participants

was determined by a questionnaire using family size (≤4, >4 persons), education status (academic or nonacademic), and house ownership (yes, no). The score of 1 was given in case of having family members of  $\leq 4$ , academic educations, and house ownership. Participants who had family members of >4, had non-academic educations, or were not house owners, were given the score of 0. The Socio-economic status score was obtained from summing up of these scores, which gave a score of 0 (poor), 1 (middle class) and 2 (high). We recorded the regular use of drugs that affect BP, including the pain medications, caffeine, and herbal supplement which used by participants then adjust the effect of drugs on BP and weight in analysis. Participants were also asked to record their daily activities during 24 hours. Mean physical activity was estimated using the following equation: PA mean =  $\Sigma$  (time activity × MET). Where, PA mean is the mean physical activity, Time activity is the total time of each activity within a day, and MET is the metabolic equivalent adapted from a reference list.34

#### Statistical analysis

All statistical analysis was performed using the IBM SPSS software version 22.0 (SPSS, Chicago, IL, USA). Normal distribution of data was checked by the Kolmogorov-Smirnov test. One-way ANOVA was used for assessed differences between tertiles of dietary GI and dietary GL. The differences between dietary intakes of participants based on the tertiles of dietary GI and dietary GL were assessed by one-way ANOVA. Logistic regression was used to assess the association of dietary GI and GL and the risk of high BP and the risk of overweight and obesity among women. Three models were constructed. Model 1 was adjusted for age and energy intake. Model 2 was adjusted for marriage status, education, occupation, number of children, family size, supplement use, medications use, and physical activity (METs/d), and Model 3 additionally adjusted for dietary intake of fiber and magnesium. Results were presented as odds ratios (ORs) and 95% confidence intervals (CIs) compared with the tertiles of dietary GI and dietary GL.

#### Results

General characteristics of participants throughout tertiles of dietary GI and dietary GL are indicated in Table 1. Totally, data on 306 women (age 32.42±8.35 years, BMI  $24.64 \pm 4.68$  kg/m<sup>2</sup>) were analyzed in the current study. Participants with the highest dietary GI tended to have lower height (P=0.01) and weight (P=0.03) than those with the lowest dietary GI. However, differences in age (P=0.36), BMI (P=0.23), waist circumference (P=0.20), SBP (P = 0.10), DBP (P = 0.22), physical activity (P = 0.16), and marriage status (P=0.52) were not significant throughout tertiles of dietary GI. In addition, subjects in the highest tertile of dietary GL had lower physical activity (P=0.05) as compared with those at the lowest tertile. Significant differences in education were found between tertiles of dietary GI and GL (both P=0.04). However, no significant differences were found in terms of age (P=0.12), weight (P=0.34), height (P=0.22), BMI (P=0.33), waist circumference (P=0.39), SBP (P=0.94), DBP (P=0.35), and marriage status (P=0.99) between

Table 1. Socio-demographic characteristics of subjects according to the tertiles of dietary glycemic index and dietary glycemic load

		Dietary glycemic index				Dietary glycemic load				
Variables	Ν	1	2	3	Р	1	2	3	<b>P</b> <sup>1</sup>	
		<59.1	59.1-61.96	>61.96	٢	<182.49	182.49-230.5	>230.5	P-	
Age (y)	306	32.75±8.19 <sup>2</sup>	33.03±8.19	31.47±8.61	0.36	32.98±7.96	33.24±7.96	31.05±8.36	0.12	
Weight (kg)	306	68.33±13.27	65.34±11.45	63.83±12.41	0.03	65.84±12.51	64.53±11.66	67.11±13.26	0.34	
Height (m)	306	163.96±5.13	163.29±5.57	161.76±5.52	0.01	163.67±5.48	163.00±5.03	162.35±5.84	0.22	
BMI (kg/m²)	306	25.29±5.01	24.31±4.32	24.32±4.67	24.32±4.67 0.23		24.23±4.18	25.18±5.10	0.33	
Waist circumference (cm)	306	87.05±12.46	84.27±11.58	84.98±10.60	0.20	85.35±11.66	84.35±9.78	86.58±13.09	0.39	
SBP (mm Hg)	306	116.72±13.28	114.12±12.91	112.82±13.55 0.10		114.90±13.48	114.32±12.62	114.45±13.90	0.94	
DBP (mm Hg)	306	73.89±6.12	72.89±6.23	72.42±6.31	72.42±6.31 0.22		73.72±5.76	72.46±6.82	0.35	
Physical activity (METs/d)	306	31.42±3.73	31.42±3.73 30.77±3.21 30.50±3.52 0.16 30		30.77±3.56	31.55±3.65	30.37±3.21	0.05		
Marriage status (%)									0.99	
Married	136	53.5%	52.5%	59.8%		55.4%	54.9%	55.4%		
Single	168	46.5%	47.5%	40.2% 0.52		44.6%	45.1%	44.6%		
Education (%)										
Low-educated	101	4.0%	17.8%	78.2%		4.0%	3.9%	10.9%	0.04	
Diploma	102	6.9%	30.7%	62.4%		20.8%	30.4%	30.7%		
Academic	103	7.8%	33.3%	58.8%	0.04	75.2%	65.7%	58.4%		

BMI, body mass index; SBP, systolic blood pressure; DBP, diastolic blood pressure.

<sup>1</sup>*P* values are resulted from ANOVA for continues variables and chi-square test for qualitative variables.

<sup>2</sup>Data are indicated as mean ± SD otherwise indicated.

#### tertiles of dietary GL.

Dietary intakes of participants based on tertiles of dietary GI and GL are shown in Table 2. Dietary intakes of total energy (P=0.006), protein (P=0.001), fat (P=0.0001), vitamin B2 (P=0.001), vitamin C (P=0.0001),  $\beta$ -carotene (P=0.0001), calcium (P=0.0001), potassium (P=0.0001), fruit (*P*=0.0001), vegetable (*P*=0.0001), dairy (*P*=0.0001) and meat (P=0.0001) were significantly lower among participants in the 3rd tertile than those in the first tertile of dietary GI. Furthermore, participants in the highest tertile of dietary GI had higher intakes of carbohydrate (P=0.0001), vitamin B1 (P=0.0001), and refined grains (P=0.0001) as compared to those in the lowest tertile. With regards to the dietary GL, lower intakes of protein (P=0.0001), vitamin B2 (P=0.0001), calcium (P=0.002), and potassium (P=0.0001) were found among subjects in the 3rd versus those in the first tertile of dietary GL. In addition, participants with the highest dietary GL had significantly higher intake of energy (P=0.0001), carbohydrate (P=0.0001), fat (P=0.002), vitamin B1 (P=0.03), whole grains (P=0.0001), refined grains (P=0.0001) and fruits (P=0.04) than those with the lowest dietary GL. Differences in intakes of vitamin C (P=0.16), vitamin E (P=0.21),  $\beta$ -carotene (P=0.26), sodium (P=0.20), vegetable (P=0.89), dairy (P=0.18) and meat (P=0.90) were not statistically significant between the highest as compared to the lowest tertile of dietary GL.

Multivariate-adjusted models with 95% confidence intervals for risk of high SBP and DBP across tertiles

of dietary GI and GL have been indicated in Table 3. In the crude model, no significant association was found between GI with SBP and DBP (for SBP: OR: 0.71; 95% CI: 0.34-1.30, 0.30; for DBP: OR: 0.61; 95% CI: 0.34-1.10, P = 0.10). In addition, no significant correlation was found between GL with SBP and DBP (for SBP: OR: 1.11; 95% CI: 0.34-1.38, P=0.73; For DBP: OR: 0.95; 95% CI: 0.53-1.72, P=0.88). Furthermore, after adjustment for the confounders including age, energy intake, marriage status, education, occupation, number of children, family size, supplement and medications use, physical activity, as well as dietary intake of fiber and magnesium in the final model, the associations remained unchanged; Such that no significant association were found between GI and SBP (OR: 0.96; 95% CI: 0.42-2.17, *P*=0.87), GI and DBP (OR: 0.72; 95% CI: 0.35-1.45, *P*=0.37), GL and SBP (OR: 1.04; 95% CI: 0.43-2.49, P=1.00), as well as between GL and DBP (OR: 1.20; 95% CI: 0.56-2.00, P=0.61).

Table 4 shows multivariate-adjusted models with 95% confidence intervals for risk of obesity and overweight across tertiles of dietary GI and GL. In the crude model before adjustment for the confounders, participants in the highest tertile of GL were at 80% higher risk for obesity and overweight (OR: 1.80; 95% CI: 1.02-3.16, P=0.03) than those in the lowest tertile. After controlling for the potential confounders, the association disappeared. In addition, differences in obesity and overweight between the highest rather the lowest tertile of GI were not significant (OR: 0.75; 95% CI: 0.42-1.31, P=0.31). This

Table 2. Dietary intakes of participants based on the tertiles of dietary glycemic index and dietary glycemic load

	Ter	tiles of dietary glyo	cemic index	Tertiles of dietary glycemic load				
Variables	1 (n=102)	2 (n=102)	3 (n=102)	Р	1 (n=102)	2 (n=102)	3 (n=102)	<b>P</b> <sup>1</sup>
Energy (kcal)	2289.17±704.10 <sup>2,3</sup>	2238.69±712.98	1998.93±622.88	0.006	1877.81±490.70	2083.84±647.46	2565.15±727.57	0.0001
Protein (g)	81.58±15.85	76.78±19.51	73.59±9.40	0.001	82.31±20.20	77.07±12.14	72.56±12.23	0.0001
Carbohydrate (g)	318.9±37.67	332.99±30.03	340.91±29.16	0.0001	316.83±30.64	330.76±29.41	345.27±34.77	0.0001
Fat (g)	70.17±14.49	64.76±15.30	61.74±12.35	0.0001	61.06±20.05	63.07±25.84	72.52±28.18	0.002
Vitamin B1 (mg)	1.84±0.36	1.91±0.38	2.06±0.03	0.0001	1.86±0.33	1.96±0.25	1.99±0.48	0.035
Vitamin B2 (mg)	2.24±0.48	2.01±0.40	1.74±0.35	0.0001	2.13±0.44	1.98±0.44	1.87±0.47	0.0001
Vitamin C (mg)	163.56±69.61	132.15±55.42	108.74±46.74	0.0001	140.01±51.78	139.22±65.21	125.21±67.50	0.160
Vitamin E (mg)	10.67±4.90	10.59±4.65	12.08±5.91	0.071	11.53±5.25	11.43±5.25	10.38±5.10	0.217
β-Carotene (µg)	907.22±1073.33	600.68±448.41	473.01±430.92	0.0001	705.82±69.88	924.51±91.54	521.82±51.66	0.268
Calcium (mg)	1130.03±309.54	1002.61±269.71	867.79±181.62	0.0001	1076.46±297.77	977.95±257.90	946.03±267.89	0.002
Potassium (mg)	3730.25±833.88	3185.97±892.58	2754.78±552.33	0.0001	932.87±92.36	785.24±77.75	774.93±76.72	0.0001
Sodium (mg)	4998.15±2869.99	5215.41±2425.15	5235.11±2150.25	0.754	5431.51±2571.15	5200.24±2481.00	4816.92±2411.61	0.206
Whole grain (g)	5132.66±79.30	5131.01±86.47	5113.03±72.47	0.150	5101.16±65.83	5119.96±62.57	5155.57±97.29	0.0001
Refined grain (g)	281.17±108.12	373.57±104.81	481.80±137.02	0.0001	274.66±89.87	388.14±89.87	473.74±89.87	0.0001
Fruit (g)	468.12±248.43	388.25±187.66	265.49±151.22	0.0001	338.05±189.14	370.39±219.74	413.42±232.15	0.043
Vegetable (g)	466.59±241.43	338.55±157.70	291.35±160.10	0.0001	366.58±200.18	371.63±212.57	358.28±200.12	0.895
Dairy (g)	614.65±277.33	529.15±270.40	371.25±168.49	0.0001	479.06±230.35	492.28±261.26	543.70±292.47	0.180
Meat (g)	147.19±85.39	123.40±55.35	108.18±50.40	0.0001	125.70±66.02	124.50±65.79	128.56±70.58	0.907

<sup>1</sup>P values are resulted from ANOVA (Analysis of variance).

<sup>2</sup> Mean ± SD.

<sup>3</sup> Data are adjusted for energy intake.

	Glycemic index				Glycemic load				
	1	2	3	Pa	1	2	3	Р	
High systolic blood pressure									
Crude	1	0.66 (0.34, 1.30)	0.71 (0.34,1.30)	0.30	1	0.68 (0.34, 1.38)	1.11 (0.34,1.38)	0.73	
Model 1 <sup>1</sup>	1	0.63 (0.31, 1.27)	0.86 (0.43,1.74)	0.63	1	0.59 (0.28, 1.24)	0.92 (0.43,1.99)	0.80	
Model 2 <sup>2</sup>	1	0.64 0.31, 1.32)	0.81 (0.37,1.73)	0.53	1	0.48 (0.22, 1.05)	0.86 (0.38,1.94)	0.65	
Model 3 <sup>3</sup>	1	0.70 (0.33, 1.48)	0.96 (0.42,2.17)	0.87	1	0.52 (0.24, 1.16)	1.04 (0.43,2.49)	1.00	
High diastolic blood pressure									
Crude	1	0.74 (0.41, 1.31)	0.61 (0.34,1.10)	0.10	1	1.19 (0.66, 2.12)	0.95 (0.53,1.72)	0.88	
Model 1	1	0.71 (0.40, 1.27)	0.61 (0.33,1.11)	0.10	1	1.19 (0.66,2.15)	1.03 (0.53, 1.97)	0.90	
Model 2	1	0.75 (0.41, 1.38)	0.69 (0.36,1.33)	0.26	1	1.03 (0.55, 1.92)	1.05 (0.52,2.09)	0.88	
Model 3	1	0.77 (0.41, 1.45)	0.72 (0.35,1.45)	0.37	1	1.06 (0.56, 2.00)	1.20 (0.56,2.00)	0.61	

<sup>a</sup> *P* values are from logistic regression.

<sup>1</sup>Model 1: Adjusted for age and energy intake.

<sup>2</sup>Model 2: Further adjusted for marriage status, education, occupation, number of children, family size, supplement use, medications use, and physical activity (METs/d).

<sup>3</sup>Model 3: Further adjusted for dietary intake of fiber and magnesium.

Table 4. Association of dietary glycemic index and glycemic load and the risk of overweight and obesity among Tehranian women.

		Glycemic index			Glycemic load				
	1	2	3	$P^{a}$	1	2	3	Р	
Crude	1	0.87 (0.49,1.51)	0.75 (0.42,1.31)	0.31	1	1.08 (0.61,1.93)	1.80 (1.02,3.16)	0.03	
Model 1 <sup>1</sup>	1	0.89 (0.47,1.66)	1.03 (0.54,1.95)	0.92	1	0.94 (0.49,1.80)	1.79 (0.89,3.62)	0.11	
Model 2 <sup>2</sup>	1	0.83 (0.43,1.60)	0.86 (0.43,1.71)	0.67	1	0.62 (0.34,1.37)	1.48 (0.70,3.16)	0.35	
Model 3 <sup>3</sup>	1	0.81(0.40,1.64)	0.83 (0.39,1.79)	0.65	1	0.70 (0.35,1.41)	1.54 (0.70,3.40)	0.34	

<sup>a</sup> *P* values are from logistic regression.

<sup>1</sup>Model 1: Adjusted for age and energy intake.

<sup>2</sup>Model 2: Further adjusted for marriage status, education, occupation, number of children, family size, supplement use, medications use, and physical activity (METs/d).

<sup>3</sup>Model 3: Further adjusted for dietary intake of fiber and magnesium.

association remained non-significant after controlling for the age, energy intake, marriage status, education, occupation, number of children, family size, supplement and medications use, physical activity, dietary intake of fiber and magnesium(OR: 0.83; 95% CI: 0.39-1.79, P=0.65). Differences in obesity and overweight between tertiles of GI were also non-significant (OR: 0.75; 95% CI: 0.42-1.31, P=0.31), even after adjustment for the potential confounders (OR: 1.54; 95% CI: 0.70-3.40, P=0.26).

#### Discussion

Findings from this cross-sectional study did not show significant association between dietary GI/GL and odds of high SBP/DBP. In addition, we could not find a significant association between dietary GI/GL and risk of obesity and overweight. Although there were differences of energy intake with GI and GL between individuals, data are adjusted for energy intake in Table 2.

These findings are in the same line with a study by Sloth et al in which differences in DBP and SBP were not significant in participants who consumed a low GI diet than those with a high GI diet.<sup>21</sup> Moreover, another study in 2010 did not find a significant association between dietary GI/GL and SBP/DBP, among 878 postmenopausal women.<sup>35</sup> On the other hand, some other studies have found a significant association between dietary GI/GL and BP in adults and aged peoples.<sup>23,36-39</sup> It should be noted that participants in these studies were from both genders and suffered from hypertension. Therefore, one may expect that the association of dietary GI/GL with BP is different among males comparing to females.

It is necessary to mention consuming a high carbohydrates diet enhances postprandial glycemia and insulin secretion at various speeds, depending on the source of carbohydrates as well as amount and type of the dietary fibers.<sup>40</sup> Therefore, quality and quantity of the ingested carbohydrates are the principal determinants of postprandial glycemic response.<sup>41</sup>

In this study, we also found that women who consumed a diet with higher GI were more likely to have higher intakes of vitamin B1, vitamin B2, vitamin C,  $\beta$ -carotene, calcium, and potassium. Also, there was a direct association between dietary GL and intakes of some nutrients including vitamin C, B1, B2, P, and calcium. Regarding to these findings, it can be suggested that those with the higher dietary GI/GL have also higher intakes of some vitamins and minerals which have found to have anti-hypertension effects.<sup>42-44</sup>

In current study, we found a direct association between dietary GL and odds of overweight and obesity, however, after adjustment for the potential confounders this association disappeared. In addition, we failed to find a significant association between dietary GI and odds of overweight and obesity, even after adjustment for the potential confounders. In line with our findings, some studies could not find differences in body weight between subjects who consumed a high or a low GI diet.<sup>21,45,46</sup> In contrast, some other studies have reported an inverse association between consumption of a low GI diet and body weight.<sup>47,48</sup> These different findings may be partially due to differences in study design, target population, and dietary assessment tools used to determine dietary GI and GL.

To the best of our knowledge, current study is the first study investigating association of dietary GI/GL with high BP among adult women in Iran. Some limitations should be kept in the mind. Due to the cross-sectional design of this study, causality could not be discovered. Hence, future researches, in particular randomized clinical trials are required to confirm these findings and to specify the causality. Although, we used a validated questionnaire to estimate dietary intakes of participants, FFQ has not been particularly planned to evaluate dietary GL and GI, therefore, it should be used carefully. Besides, due to using FFQ, misclassification of study participants is also probable.

In conclusion, we could not find significant association between dietary GI/GL and the risk of high SBP/DBP, even after controlling for a wide range of potential confounding factors. In addition, no association was found between dietary GI/GL and odds of overweight or obesity in adult women. Further studies, in particular large scale clinical trials are required to shed light in this area.

#### **Competing interests**

The authors have no conflicts of interest to declare and they did not use any outside assistance in preparing the manuscript.

#### **Ethical approval**

The study protocol was approved by the local ethics committee of the Tehran University of Medical Sciences (Code: BN092).

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#### References

1. Carr DB, Utzschneider KM, Hull RL, Kodama K, Retzlaff

BM, Brunzell JD, et al. Intra-abdominal fat is a major determinant of the National Cholesterol Education Program Adult Treatment Panel III criteria for the metabolic syndrome. **Diabetes** 2004;53(8):2087-94. doi: 10.2337/diabetes.53.8.2087

- Ren Q, Su C, Wang H, Wang Z, Du W, Zhang B. Prospective Study of optimal obesity index cut-off values for predicting incidence of hypertension in 18-65-year-old chinese adults. PloS One. 2016;11(3):e0148140. doi: 10.1371/journal. pone.0148140.
- Kingue S, Ngoe CN, Menanga AP, Jingi AM, Noubiap JJ, Fesuh B, et al. Prevalence and risk factors of hypertension in urban areas of cameroon: a nationwide population-based cross-sectional study. J Clin Hypertens 2015;17(10):819-24. doi: 10.1111/jch.12604.
- 4. Esteghamati A, Abbasi M, Alikhani S, Gouya MM, Delavari A, Shishehbor MH, et al. Prevalence, awareness, treatment, and risk factors associated with hypertension in the Iranian population: the national survey of risk factors for noncommunicable diseases of Iran. **Am J Hypertens** 2008;21(6):620-6. doi: 10.1038/ajh.2008.154
- Noohi F, Sarrafzadegan N, Khosravi A, Andalib E. The first Iranian recommendations on prevention, evaluation and management of high blood pressure. ARYA Atheroscler 2012;8(3):97-118.
- Seo MH, Lee J-Y, Ryu S, Won YS, Sung KC. The Effects of urinary albumin and hypertension on all-cause and cardiovascular disease mortality in Korea. Am J Hypertens 2017;30(8):799-807. doi: 10.1093/ajh/hpx051.
- Wang L, Manson JE, Gaziano JM, Liu S, Cochrane B, Cook NR, et al. Circulating inflammatory and endothelial markers and risk of hypertension in white and black postmenopausal women. Clin Chem 2011;57(5):729-36. doi: 10.1373/clinchem.2010.156794.
- Defagó MD, Gu D, Hixson JE, Shimmin LC, Rice TK, Gu CC, et al. Common genetic variants in the endothelial system predict blood pressure response to sodium intake: the GenSalt study. Am J Hypertens 2013;26(5):643-56. doi: 10.1093/ajh/hps099.
- Brown IJ, Stamler J, Van Horn L, Robertson CE, Chan Q, Dyer AR, et al. Sugar-sweetened beverage, sugar intake of individuals, and their blood pressure international study of macro/micronutrients and blood pressure. Hypertension 2011;57(4):695-701. doi: 10.1161/ HYPERTENSIONAHA.110.165456.
- Hosseininasab M, Norouzy A, Nematy M, Bonakdaran S. Low-glycemic-index foods can decrease systolic and diastolic blood pressure in the short term. Int J Hypertens 2015;2015:801268. doi: 10.1155/2015/801268.
- Gopinath B, Flood VM, Rochtchina E, Baur LA, Smith W, Mitchell P. Influence of high glycemic index and glycemic load diets on blood pressure during adolescence. Hypertension. 2012;59(6):1272-7. doi: doi: 10.1161/ HYPERTENSIONAHA.112.190991.
- Carels RA, Blumenthal JA, Sherwood A. Emotional responsivity during daily life: Relationship to psychosocial functioning and ambulatory blood pressure. Int J Psychophysiol. 2000;36(1):25-33.
- Lima ST, de Souza BD, França AK, Salgado Filho N, Sichieri R. Dietary approach to hypertension based on low glycaemic index and principles of DASH (Dietary

Approaches to Stop Hypertension): a randomised trial in a primary care service. **Br J Nutr** 2013;110(8):1472-9. doi: 10.1017/S0007114513000718.

- Shah M, Adams-Huet B, Garg A. Effect of highcarbohydrate or high-cis-monounsaturated fat diets on blood pressure: a meta-analysis of intervention trials. Am J Clin Nutr 2007;85(5):1251-6. doi: 10.1093/ajcn/85.5.1251.
- Rouhani MH, Kelishadi R, Hashemipour M, Esmaillzadeh A, Azadbakht L. The effect of low glycemic index diet on body weight status and blood pressure in overweight adolescent girls: a randomized clinical trial. Nutr Res Pract 2013;7(5):385-92. doi: 10.4162/nrp.2013.7.5.385.
- Hosseininasab M, Norouzy A, Nematy M, Bonakdaran S. Low-glycemic-index foods can decrease systolic and diastolic blood pressure in the short term. Int J Hypertens 2015;2015:801268. doi: 10.1155/2015/801268.
- Lima ST, Souza BS, Franca AK, Salgado JV, Salgado-Filho N, Sichieri R. Reductions in glycemic and lipid profiles in hypertensive patients undergoing the Brazilian Dietary Approach to Break Hypertension: a randomized clinical trial. Nutr Res 2014;34(8):682-7. doi: 10.1016/j. nutres.2014.07.009.
- Evans CE, Greenwood DC, Threapleton DE, Gale CP, Cleghorn CL, Burley VJ. Glycemic index, glycemic load, and blood pressure: a systematic review and metaanalysis of randomized controlled trials. Am J Clin Nutr 2017;105(5):1176-90. doi: 10.3945/ajcn.116.143685.
- Castro-Quezada I, Artacho R, Molina-Montes E, Serrano FA, Ruiz-Lopez MD. Dietary glycaemic index and glycaemic load in a rural elderly population (60-74 years of age) and their relationship with cardiovascular risk factors. Eur J Nutr 2015;54(4):523-34. doi: 10.1007/s00394-014-0733-9.
- Heidari-Beni M, Golshahi J, Esmaillzadeh A, Azadbakht L. Potato consumption as high glycemic index food, blood pressure, and body mass index among Iranian adolescent girls. ARYA Atheroscler 2015;11(Suppl 1):81-7.
- Sloth B, Krog-Mikkelsen I, Flint A, Tetens I, Björck I, Vinoy S, et al. No difference in body weight decrease between a low-glycemic-index and a high-glycemic-index diet but reduced LDL cholesterol after 10-wk ad libitum intake of the low-glycemic-index diet. Am J Jlin Nut. 2004;80(2):337-47. doi: 10.1093/ajcn/80.2.337
- 22. Flight I, Clifton P. Cereal grains and legumes in the prevention of coronary heart disease and stroke: a review of the literature. **Eur J Clin Nutr** 2006;60(10):1145-59. doi:10.1038/sj.ejcn.1602435.
- Castro-Quezada I, Artacho R, Molina-Montes E, Serrano FA, Ruiz-López MD. Dietary glycaemic index and glycaemic load in a rural elderly population (60–74 years of age) and their relationship with cardiovascular risk factors. Eur J Clin Nutr 2015;54(4):523-34. doi: 10.1007/s00394-014-0733-9.
- Juanola-Falgarona M, Salas-Salvadó J, Buil-Cosiales P, Corella D, Estruch R, Ros E, et al. Dietary glycemic index and glycemic load are positively associated with risk of developing metabolic syndrome in middle-aged and elderly adults. J Am Geriatr Soc 2015;63(10):1991-2000. doi: 10.1111/jgs.13668.
- 25. Hosseinpour-Niazi S, Sohrab G, Asghari G, Mirmiran P, Moslehi N, Azizi F. Dietary glycemic index, glycemic load, and cardiovascular disease risk factors: Tehran Lipid and

Glucose Study. Arch Iran Med. 2013;16(7):401-7.

- 26. Azadbakht L, Esmaillzadeh A. Red meat intake is associated with metabolic syndrome and the plasma C-reactive protein concentration in women. **J Nutr** 2008;139(2):335-9. doi: 10.3945/jn.108.096297.
- Haghighatdoost F, Azadbakht L, Keshteli AH, Feinle-Bisset C, Daghaghzadeh H, Afshar H, et al. Glycemic index, glycemic load, and common psychological disorders. Am J Clin Nutr 2016;103(1):201-9. doi: 10.3945/ajcn.114.105445.
- 28. Taleban F, Esmaeili M. Glycemic index of Iranian foods: Guideline for diabetic and hyperlipidemic patients. Tehran: National Nutrition and Food Technology of Iran, Shahid Beheshti University of Medical Science; 1999.
- Atkinson FS, Foster-Powell K, Brand-Miller JC. International tables of glycemic index and glycemic load values: 2008. Diabetes Care 2008;31(12):2281-3. doi: 10.2337/dc08-1239.
- Foster-Powell K, Holt SH, Brand-Miller JC. International table of glycemic index and glycemic load values: 2002. Am J Clin Nutr 2002;76(1):5-56. doi:10.1093/ajcn/76.1.5.
- Brand-Miller J, Foster-Powell K, Colagiuri S, Barclay A. The New Glucose Revolution for Diabetes: The Definitive Guide to Managing Diabetes and Prediabetes Using the Glycemic Index. Hachette UK: Da Capo Lifelong Books; 2007.
- Wolever TM, Yang M, Zeng XY, Atkinson F, Brand-Miller JC. Food glycemic index, as given in glycemic index tables, is a significant determinant of glycemic responses elicited by composite breakfast meals. Am J Clin Nutr 2006;83(6):1306-12. doi:10.1093/ajcn/83.6.1306.
- 33. Izadi V, Esmaillzadeh A, Hashemipour M, Surkan PJ, Azadbakht L, Kelishadi R. High protein diets do not affect anthropometric indexes and cardiometabolic risk factors among children with excess weight: A randomized controlled trial. J Cardiovasc Thorac Res 2018;10(2):95. doi: 10.15171/jcvtr.2018.15.
- 34. Ainsworth BE, Haskell WL, Whitt MC, Irwin ML, Swartz AM, Strath SJ, et al. Compendium of physical activities: an update of activity codes and MET intensities. Med Sci Sports Exerc 2000;32(9 suppl):S498-504.
- 35. Shikany JM, Tinker LF, Neuhouser ML, Ma Y, Patterson RE, Phillips LS, et al. Association of glycemic load with cardiovascular disease risk factors: the Women's Health Initiative Observational Study. Nutrition 2010;26(6):641-7. doi: 10.1016/j.nut.2009.08.014.
- 36. Lin P-H, Chen C, Young DR, Mitchell D, Elmer P, Wang Y, et al. Glycemic index and glycemic load are associated with some cardiovascular risk factors among the PREMIER study participants. Food Nutr Res 2012;56(1):9464. doi: 10.3402/fnr.v56i0.9464.
- 37. Milton JE, Briche B, Brown IJ, Hickson M, Robertson CE, Frost GS. Relationship of glycaemic index with cardiovascular risk factors: analysis of the National Diet and Nutrition Survey for people aged 65 and older. Public Health Nutr 2007;10(11):1321-35. doi: 10.1017/S1368980007702914.
- 38. Wolever TM, Gibbs AL, Mehling C, Chiasson J-L, Connelly PW, Josse RG, et al. The Canadian Trial of Carbohydrates in Diabetes (CCD), a 1-y controlled trial of low-glycemicindex dietary carbohydrate in type 2 diabetes: no effect on glycated hemoglobin but reduction in C-reactive protein. Am J Clin Nutr 2008;87(1):114-25. doi: 10.1093/

ajcn/87.1.114.

- Maki KC, Rains TM, Kaden VN, Raneri KR, Davidson MH. Effects of a reduced-glycemic-load diet on body weight, body composition, and cardiovascular disease risk markers in overweight and obese adults. Am J Clin Nutr 2007;85(3):724-34. doi:10.1093/ajcn/85.3.724.
- Turati F, Galeone C, Gandini S, Augustin LS, Jenkins DJ, Pelucchi C, et al. High glycemic index and glycemic load are associated with moderately increased cancer risk. Mol Nutr Food Res 2015;59(7):1384-94. doi: 10.1002/ mnfr.201400594.
- 41. Howarth NC, Saltzman E, Roberts SB. Dietary fiber and weight regulation. **Nutr Rev** 2001;59(5):129-39.
- 42. Houston MC. The role of cellular micronutrient analysis, nutraceuticals, vitamins, antioxidants and minerals in the prevention and treatment of hypertension and cardiovascular disease. Ther Adv Cardiovasc Dis 2010;4(3):165-83. doi: 10.1177/1753944710368205.
- Taddei S, Virdis A, Ghiadoni L, Magagna A, Salvetti A. Vitamin C improves endothelium-dependent vasodilation by restoring nitric oxide activity in essential hypertension. Circulation. 1998;97(22):2222-9.
- 44. Egeland GM, Skurtveit S, Sakshaug S, Daltveit AK, Vikse BE, Haugen M. Low Calcium Intake in Midpregnancy Is

Associated with Hypertension Development within 10 Years after Pregnancy: The Norwegian Mother and Child Cohort Study. **J Nutr** 2017;12(10):251520. doi: 10.3945/jn.117.251520.

- Pereira EV, Costa Jde A, Alfenas Rde C. Effect of glycemic index on obesity control. Arch Endocrinol Metab 2015;59(3):245-51. doi: 10.1590/2359-3997000000045.
- 46. de la Fuente-Arrillaga C, Martinez-Gonzalez MA, Zazpe I, Vazquez-Ruiz Z, Benito-Corchon S, Bes-Rastrollo M. Glycemic load, glycemic index, bread and incidence of overweight/obesity in a Mediterranean cohort: the SUN project. BMC Public Health 2014;14:1091. doi: 10.1186/1471-2458-14-1091.
- Feliciano Pereira P, das Gracas de Almeida C, Alfenas Rde C. Glycemic index role on visceral obesity, subclinical inflammation and associated chronic diseases. Nutr Hosp 2014;30(2):237-43. doi: 10.3305/nh.2014.30.2.7506.
- Schwingshackl L, Hoffmann G. Long-term effects of low glycemic index/load vs. high glycemic index/load diets on parameters of obesity and obesity-associated risks: a systematic review and meta-analysis. Nutr Metab Cardiovasc Dis 2013:23(8):699-706. doi: 10.1016/j. numecd.2013.04.008.

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